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- 7) Applicant: KYOWA HAKKO KOGYO KABUSHIKI KAISHA 6-1, Ohte-machi 1-chome Chiyoda-ku Tokyo 100(JP)
- 7) Applicant: YAMAHATSU, SANGYO CO., LTD. 1-1-25, Dojima, Kita-ku Osaka-shi Osaka 530(JP)
- (72) Inventor: TSUJINO, Yoshio 1-1-31, Hagurazaki Izumisano-shi Osaka 598(JP)
- (72) Inventor: YOKOO, Yoshiharu 3-4-17, Yokoyama Sagamihara-shi Kanagawa 229(JP)
- (72) Inventor: SAKATO, Kuniaki 987-37, Kamiogino Atsugi-shi Kanagawa 243-02(JP)
- (72) Inventor: HAGINO, Hiroshi 3-16-1-701, Kaminoge Setagaya-ku Tokyo 158(JP)
- (4) Representative: Vossius & Partner Siebertstrasse 4 P.O. Box 86 07 67 D-8000 München 86(DE)

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54) HAIR-CARE PRODUCT COMPOSITION.

The present invention provides a hair preparation composition which contains, as an active ingredient, a dielectron reducing oxidase utilizing oxygen as an acceptor.

Croydon Printing Company Ltd.

#### SPECIFICATION

#### HAIR PREPARATION COMPOSITION

## Technical Field

The present invention relates to a hair preparation composition characterized in that a dielectron reducing oxidase utilizing oxygen as an acceptor is incorporated as an active ingredient.

#### Background Art

Heretofore, with oxidation hair dyes, for oxidation polymerization of an oxidation dye and in order to bleach hair, hydrogen peroxide, sodium perborate, sodium percarbonate, etc. are used as the oxidizing agent.

Further, in general, for performing a permanent wave, there is employed a method which comprises reducingly cutting the S-S bond in hair with a first solution containing a reducing agent such as thioglycolic acid, cysteine, etc. as a main component and thereafter oxidizingly fixing this hair with a second solution containing an oxidizing agent such as sodium bromate, sodium perborate, hydrogen peroxide etc. as a main component.

However, the use of these oxidizing agents has a disadvantage that they are apt to damage hair and skin.

#### Disclosure of the Invention

The present invention is characterized by activating oxygen in air with an oxidase and effectively employing its oxidizing power in the oxidizing course needed for the product.

A first object of the present invention is to provide a hair preparation composition which has a satisfactory effect with a mild oxidizing effect.

A second object of the present invention is to

provide a hair preparation composition having an oxidizing function which is low in skin irritation and hardly damages hair and skin, such as an oxidation hair dye, an oxidation agent composition for permanent waving preparation, a bleaching agent for hair or body hair on hands and feet, etc.

The present invention is described in more detail below.

The present invention provides a hair preparation composition which contains at least one dielectron reducing oxidase utilizing oxygen as an acceptor.

The enzyme used in the present invention is a dielectron reducing oxidase utilizing oxygen as an acceptor, and mention is made of, for example, pyranose oxidase (hereinafter referred to as PROD), glucose oxidase (hereinafter referred to as GOD), glycerol oxidase, lactic acid oxidase, pyruvic acid oxidase, uricase, etc.

In the present invention, these enzymes are used either alone or as a combination of two or more thereof.

Further, in order that certain enzymes suitable for the present invention function, cofactors as required. For example, it is known that as cofactors, flavin adenine dinucleotide (hereinafter referred to as FAD) and iron ions are necessary for PROD; FAD and iron ions for GOD; and FAD for pyruvic acid oxidase.

These cofactors are generally contained as impurities in enzymes, and therefore, where the enzyme is described in the present specification, it is understood that it also contains the necessary cofactors unless otherwise specified.

The present invention further provides a hair preparation composition which contains the above-described oxidase and a donor of said enzyme.

The donor varies depending on the enzyme, and for example, there are used D-glucose, L-sorbose and D-xylose for PROD; D-glucose for GOD; glycerol, dihydroxyacetone for

glycerol oxidase; lactic acid and its salts for lactic acid oxidase; pyruvic acid and its salts for pyruvic acid oxidase; and uric acid and its salts for uricase.

The amount of the enzyme incorporated is 1 unit/100g to 1 x  $10^8$  units/100g, preferably 1 x  $10^2$  units/100g to 1 x  $10^5$  units/100g, as the concentration in actual use on hair.

The amount of the donor is 0.01% by weight to 60% by weight, preferably 1% by weight to 35% by weight, as the concentration in actual use on hair.

In the present invention, where D-glucose is used as the donor, PROD is preferred as the enzyme. This is because PROD acts on both  $\alpha$ -type and  $\beta$ -type of D-glucose.

Where GOD is used, it is preferred to use it in combination with mutarotase. This is because mutarotase performes rearrangement of D-glucose from  $\alpha$ -type to  $\beta$ -type.

It is known that the activity of enzymes can be maintained and stabilized at as high level as possible by the enzyme immobilization method. The immobilized enzyme can be used in the present invention.

In order to solubilize the enzyme also in an organic solvent and further in order that the activity of the enzyme may manifest itself in an organic solvent, it is possible to chemically bind a synthetic polymer or a natural polymer onto the surface of the enzyme protein molecules. This modified enzyme is also included in the enzymes of the present invention.

The present invention also provides a hair dye composition which contains an oxidation hair dye, the above-described enzyme and/or a donor of said enzyme.

The hair dye composition of the present invention is employed in three forms, namely, (1) a hair dye consisting of three packages, (2) a hair dye consisting of two packages, and (3) a hair dye consisting of one package.

As the dye used for the oxidation hair dye, all of the ordinary oxidation dyes may be used. In addition to

resorcin, there are described in Hair Dye Standard Material Specifications (third revision, May 1985, Japan Hair Color Industrial Society, Hair Dye Meeting), for example, 5aminoorthocresol, 3,3'-iminodiphenol, 2,4-diaminophenol hydrochloride, toluene-2,5-diamine hydrochloride, paraphenylenediamine hydrochloride, Nphenylparaphenylenediamine hydrochloride, metaphenylenediamine hydrochloride, orthoaminophenol, catechol, Nphenylparaphenylenediamine acetate, 2,6-diaminopyridine, 1,5dihydroxynaphthalene, diphenylamine, toluene-2,5-diamine, toluene-3,4-diamine, α-naphthol, paraaminophenylsulfamic acid, paraaminophenol, paraphenylenediamine, paramethylaminophenol, hydroquinone, pyrogallol, N-phenylparaphenylenediamine, phloroglucine, metaaminophenol, metaphenylenediamine, 5aminoorthocresol sulfate, orthoaminophenol sulfate, orthochloroparaphenylenediamine sulfate, 4,4'diaminodiphenylamine sulfate, toluene-2,5-diamine sulfate, paraaminophenol sulfate, paraphenylenediamine sulfate, paramethylaminophenol sulfate, metaaminophenol sulfate and metaphenylenediamine sulfate.

Further, 2,4-diaminophenoxyethanol hydrochloride and 5-(2-hydroxyethylamino)-2-methylphenol were also added.

Furthermore, in general, direct dyes often used in combination with oxidation dyes are also included in the oxidation dyes in a wide sense, for example, 2-amino-4-nitrophenol, 2-amino-5-nitrophenol, 1-amino-4-nethylaminoanthraquinone, nitroparaphenylenediamine hydrochloride, 1,4-diaminoanthraquinone, nitroparaphenylenediamine, picramic acid, sodium picramate, picric acid, 2-amino-5-nitrophenol sulfate, nitroparaphenylenediamine sulfate, paranitroorthophenylenediamine sulfate, paranitrometaphenylenediamine sulfate, etc.

In general, with the oxidation hair dyes, for oxidation polymerization of the oxidation dye and in order to

bleach hair, hydrogen peroxide is used at a concentration in use of 1.5% by weight to 4.0% by weight. At this concentration, damage of hair to some degree is inevitable and also skin trouble might be caused depending on the user.

In the present invention, since oxygen in air is activated and utilized, hair damage and skin trouble are less occurred and also the same dyeing and bleaching effects as that by the conventional method may be imparted.

On the other hand, heretofore, there have been found two or three techniques which utilize an enzyme in the hair treating techniques with oxidation hair dyes.

For example, in Japanese Published Unexamined Patent Application No. 10400/1972, there is proposed a method for dyeing hair which comprises (1) a peroxidase enzyme, (2) hydrogen peroxide and (3) an oxidation dye.

Further, in Japanese Published Examined Patent Application No. 47778/1976, there is proposed a composition of dithiothreitol or/and dithioerythritol as a first solution, one, or two or more of the compounds selected from the group (1) tyrosine and/or  $DL-\beta-(3,4-dihydroxyphenyl)$  alanine or a derivative thereof, (2) 5-hydroxyindole, methyl 5,6-dihydroxyindole-2-carboxylate, tyramine, dopamine, 6-hydroxydopamine and pyrocatechol as a second solution, and a persulfuric acid salt as a third solution, which further incorporates tyrosinase in the second solution or the third solution.

Further, in Japanese Published Examined Patent Application No. 31325/1983, there is also proposed a hair dye consisting of four liquids of (1) pyrocatechol, (2) a water-soluble salt of zinc, copper or iron, (3) a peroxidase and (4) hydrogen peroxide.

These are all those utilizing only the oxidizing effect of the initially incorporated hydrogen peroxide or persulfuric acid salt, and the incorporated hydrogen peroxide or persulfuric acid salt, and the incorporated enzyme is also

different, and further, in Japanese Published Examined Patent Application Nos. 47778/1976 and 31325/1983, the precursors of the dyes are specified and thus their intentions are different from the present invention.

Furthermore, the present invention may also be applied to a bleaching agent for unwanted body hair, and a bleaching agent for hair.

In addition, the present invention provides a permanent wave oxidation composition containing a dielectron reducing oxidase utilizing oxygen as an acceptor and a donor of said enzyme, the so-called second solution. Heretofore, the first solution is a reducing agent incorporating thioglycolic acid and/or cysteine, etc., and the second solution is an aqueous solution of an oxidizing agent such as sodium bromate, potassium bromate, sodium perborate, hydrogen peroxide, etc.

In general, the concentration of the hydrogen peroxide is used at 1.0% by weight to 2.5% by weight. At this concentration, damage of hair to some degree is inevitable and skin trouble might be caused depending on the user.

In the present invention, since oxygen in air is utilized, damage of hair and skin trouble are less occurred and also the same oxidation fixing effect as that by the conventional method may be imparted.

Test Examples are given below.

The percent is % by weight, and for the units of the enzyme amounts, there were used D-glucose units for GOD and PROD; uric acid units for uricase; hydrogen peroxide units for peroxidase; and  $\alpha$ -D-glucose units for mutarotase.

### Test Example 1 Dyeing Test

As shown in Table 1, oxidizing solutions were prepared from three kinds of oxidases, i.e. GOD, PROD and uricase and also from combinations of these oxidases with mutarotase and/or peroxidase, and as comparisons, from

hydrogen peroxide and purified water alone containing no oxidizing agent.

These were tested for two items. Namely, the hair dyeing effect and the finish of hair were compared and judged according to the following methods and the results are set forth in Table 1.

#### [Testing Method]

Ten ml of a sclution obtained by mixing a dye solution and an oxidizing solution at a ratio of 1:1 is coated on a goat hair bundle of 2g in weight and 10 cm in length, treated for 30 minutes, washed with water, shampooed and dried.

The evaluation of the dyeing effect and the finish of hair were done according to the following standard.

- A: Evaluation of the dyeing effect Dyeing properties
  - ⊚ : Dyed in a thick dark brown color
    - o: Dyed in a dark brown color
    - $\Delta$ : Dyed in a shallow dark brown color
    - x : Hardly dyed
- B: Finish of hair (as compared with the untreated hair)
  - o: Soft and combing is smooth
  - $\Delta$ : Somewhat soft but combing is inferior
  - x : Softness is lost and also combing is inferior

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Finish of Hair	Dyeing Effect	Purified water	Peroxidase (112 units/mg)	Mutarotase (0.95 unit/mg)	Uricase (2.8 units/mg)	PROD (1 unit/mg)	GOD (2.4 units/mg)	Hydrogen peroxide solution (35%)	(Oxidizing Solution)	Purified water	Potassium hydroxide	Ammonium hydroxide (30%)	Thioglycolic acid	Uric acid	D-Glucose	Paraphanylenodiamino	(Dye Solution)	
0	▷	Î	 	1		E se	8.3	ı		Î	adjusted to pH 7.0		0.05	1	3.6	0.13		1-1*
0	0			0.2		1	0.3				an]usted to ph 7.0		0.05	1	3.6	0.1%		1-2*
0	0		0.0058	0.2		1	8,3				adjusted to pH 7.0		0.05	1	3.6	0.18		1-3*
0	0					20.0		l			adjusted to pH 7.0		0.05	ı	3.6	0.1%		1-4*
0	0		0.0058	1	1	20.0	1				adjusted to pH 7.0		0.05	1	3.6	0.1%		1-5*
0	0	- Balance			2.0					- Balance	adjusted to pil 7.0		0.05	3.4	1	0.1%		1-6*
0	0				2.0			-			adjusted to pH 0.0		0.05	3.4	ı	0.1%		1-7*
0	0			1	2.0	1		[			adjusted to pH 9.0		0.05	3.4		0.1%		1-8*
0	0		1	1	2.0	1	i				adjusted to pH 10.0	-	0.05	3.4		0.1%		1-9*
0	0		0.0058		2.0	1					adjusted to pH 7.0	1	0.05	3.4		0.1%		1-10*
D	×				1	!		1			adjusted to pH 7.0	<u> </u>	0.05			0.18		J-11
×	0					1		14.7			1	adjusted to pH 10.0	0.05			0.18	i i	1-12
×	0							14.7		:		: :	0.05	1 :		0.1%	:	1-1.1

As seen from the above-described results, the method utilizing hydrogen peroxide as the oxidizing agent is excellent in the dyeing properties, but poor in the finish of hair. By the method utilizing purified water alone, the hair was hardly dyed. According to the present invention, a good finish of hair can be obtained while retaining almost the same dyeing effect as that by using hydrogen peroxide as the oxidizing agent.

Further, the dyeing properties were improved by the combined use with mutarotase and/or peroxidase rather than by using GOD alone.

#### Test Example 2 Waving Test

After treating with a first solution (reducing agent) of a permanent wave formulation, there was used as a second solution (oxidation agent) a composition incorporating uricase or, as a comparison, a composition incorporating sodium bromate or a composition composed of purified water alone. Three items, i.e. the wave index, wave retention coefficient and degree of skin roughness by an interdigital dropping method were examined by the following methods and the results are set forth in Table 2.

Table 2

[First Solution (Reducing Agammonium thioglycolate (50% as thioglycolic acammonium hydroxide (30% Purified water	cid)	A	djusted d	13.0% to pH 9.0 Balance
[Second Solution (Oxidation	Agent)]			
•	2-1*	2-2*	2-3	2-4
Sodium bromate	-	-	-	8.0%
Potassium urate	2.5%	2.5%	-	-
Uricase				
(2.8 units/mg)	2.0	2.0	-	-
Peroxidase				
(ll2 units/mg)	-	0.003	-	-
Purified water		Bala	ince	
Wave index (%)	61.0	59.8	60.2	59.3
Wave retention				
coefficient (%)	58.7	62.0	4.5	62.5
Degree of skin				
roughness	0	О	0	x

<sup>\*</sup> Present invention

Note: The uricase in the formulation is added just before the treatment with the second solution.

[Wave Index and Wave Retention Coefficient Measuring Tests]

(I) Twenty hairs described below were made into a bundle, fixed to a plastic measuring comb by winding the hairs in zigzags, dipped in the first solution (reducing agent) havig the composition shown in Table 2 at 30°C for 10 minutes, then rinsed thoroughly with water, and thereafter dipped in the second solution (oxidation agent) at 30°C for 10 minutes respectively. After rinsing with water, it was removed from

the measuring comb, and the degree of waving was measured according to Kirby method (Proceedings of the Scientific Section, Vol. 26, p. 12, 1956).

Hair: Human hair of 15 cm in length (10-year-old female, untreated hair), washed with a 10% aqueous solution of sodium laurylsulfate and dried in air.

(II) The hair used in (I) was dipped in a 10% aqueous solution of sodium laurylsulfate at 60°C for 20 minutes, washed with water and maltreated. Its wave index was compared with the wave index before the maltreatment and the wave retention coefficient was calculated according to the following equation.

Wave Retention Coefficient (%) =

100 - (Wave Index - Wave Index before Maltreatment) x 100
Wave Index

[Test by an Interdigital Dropping Method]

Three parts between fingers (between the second and third fingers, between the third and fourth fingers and between the fourth and fifth fingers) of either of right and left hand of the panel were examined to confirm that there is no abnormality of the skin. Thereafter, the second solution (oxidation agent) of the composition shown in Table 2 was dropped on the interdigital parts at a rate of 0.7 ml/min by a constant flow rate pump for 10 minutes, left for 5 minutes, washed with warm water of about 40°C for 15 seconds, and dried with towel; this operation was repeated twice a day for 6 days, between the former three days and the latter three days, a two-day interval was taken, and the degree of skin roughness was observed by comparing with the hand not tested.

The experimental panel was consisted of 20 females of 22 - 52 years old. The evaluation of the degree of skin roughness was done according to the following standard.

A: Evaluation of Skin Roughness

- o: Skin roughness is not observed.
- Δ: Minute wrinkles are observed.
- x: Wrinkles, dryness and roughness of skin are observed.

From the above-described results, it is obvious that the present invention has the same waving effect as the conventional product incorporating sodium bromate and that there is almost no waving effect with purified water alone.

Further, with the product incorporating sodium bromate, skin roughness is observed, while with the oxidation agent of the present invention, the skin roughness of the panel is remarkably prevented.

#### Brief Description of the Drawing

Fig. 1 is a cross-sectional view of a piston can used in Examples of the present invention.

#### Examples

The present invention is described in more detail by the following examples.

Example 1	One-package	Hair	Dye	(Cream)
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Paraphenylenediamine	1.35%
Orthoaminophenol	0.1
Resorcin	0.25
Cetanol	6.0
Oleyl alcohol	5.0
Polyoxyethylene cetyl ether (15 E.O.)	7.0
Liquid paraffin	10.0
Stearyltrimethylammonium chloride	1.0
Propylene glycol	2.0
Uricase (2.8 units/mg)	1.0
Uric acid	5.3
Disodium edetate	0.2
Thioglycolic acid	0.1

Monoethanolamine, purified water

Balance

(The pH is adjusted to 8.5 with monoethanolamine.)

This composition is coated on a white hair, which is treated at  $30\,^{\circ}\text{C}$  for 30 minutes, then washed with water, shampooed and dried.

The white hair was dyed in grayish color.

Example 2	One-package	Hair	Dye	(Gel)
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Paraphenylenediamine	0.08%
Orthoaminophenol	0.04
Nitroparaphenylenediamine	0.4
2-Amino-4-nitrophenol	0.4
Resorcin	0.1
Sodium carboxymethyl cellolose	7.5
<pre>Uricase (2.8 units/mg)</pre>	1.07
Potassium urate	2.44
Thioglycolic acid	0.1
Purified water	Balance

This composition is coated on a white hair, which is treated at  $30\,^{\circ}\text{C}$  for 30 minutes, then washed with water, shampooed and dried.

The white hair was dyed in reddish brown color.

Example 3 One-package Hair Dye (Hair cream type)

Paraphenylenediamine		0.135%
Orthoaminophenol	•	0.01
Resorcin		0.025
Cetanol		6.0
Oleyl alcohol		5.0
Polyoxyethylene cetyl et	ther (15 E.O.)	7.0
. ·	(10 E.O.)	3.5
Liquid paraffin		10.0-
Stearyltrimethylammonium	m chloride	1.0
Liquid lanolin		1.0
<pre>Uricase (2.8 units/mg)</pre>		1.0

Uric acid	1.1%
Disodium edetate	0.2
Ascorbic acid	0.2
Potassium hydroxide, purified water	Balance
(The pH is adjusted to 7.5 with potassium	
hydroxide.)	

This composition is coated on a white hair, which is treated for 10 to 30 minutes. The hair is immediately washed or shampooed and dried.

This composition was similarly used on the white hair every day for 10 days, to find that the hair had been dyed in grayish color.

Example 4	One-package	Hair Dye	(Treatment	type)
-----------	-------------	----------	------------	-------

Paraphenylenediamine	- 0.04%
Orthoaminophenol	0.02
Nitroparaphenylenediamine	0.20
2-Amino-4-nitrophenol	0.20
Resorcin	0.05
Cetyltrimethylammonium chloride	2.5
Stearyltrimethylammonium chloride	1.0
Isopropyl myristylate	7.0
Cetanol	5.0
Stearyl alcohol	2.0
Liquid paraffin	4.0
Liquid lanolin	0.5
Propylene glycol	0.5
Uricase (2.8 units/mg)	1.0
Uric acid	5.1
Peroxidase (112 units/mg)	0.003
Thioglycolic acid	0.1
Potassium hydroxide, purified water	Balance

(The pH is adjusted to 7.5 with potassium hydroxide.)

This composition is coated on a white hair, which is treated at 30°C for 30 minutes, then washed with water,

shampooed and dried. The white hair was dyed in reddish brown color. Further, like the conventional hair treatment, it was coated on the previously shampooed hair, which was treated at 30 to  $40^{\circ}$ C for 5 to 10 minutes, and washed with water.

This precedure was repeated on the white hair for 10 days, to find that the hair was dyed in reddish brown color.

Example	5	One-package	Hair	Dye	(Powder)
	-	0 paoag		2/-	(10,000)

Paraphenylenediamine sulfate	10.0 %
Orthoaminophenol sulfate	3.0
Paramethylaminophenol sulfate	2.0
Sodium carboxymethyl cellulose	24.0
Sodium carbonate	7.36
Uric acid	14.17
<pre>Uricase (2.8 units/mg)</pre>	8.30
Dextrin	Balance

First, to 6 g of the above-described composition is added purified water until the total volume is  $50\ ml$ , for preparation of a dye solution.

This dye solution is coated on a white hair, which is treated at 30°C for 30 minutes, washed with water, shampooed and dried.

The white hair was dyed in dark reddish brown color.

In this powder hair dye, where water is not contained in the starting material, the stability was improved.

Example 6 Aerosol Type Hair Dye [Stock Solution]

Paraphenylenediamine	0.4%
Paraaminophenol	0.1
Orthoaminophenol	0.5
Resorcin	0.8
PROD (1 unit/mg)	10.0
D-Glucose	5.5
Propylene glycol	2.0

## Thioglycolic acid

0.1%

#### 0.1M Phosphate buffer (pH 6.5)

Balance

First, the above-described composition is injected into a piston can (double container system, Fig. 1), and then carbon dioxide is injected from a gas filling line so that the initial volume be about 1/3 of the container and the initial pressure be about 8 to 9 kg/cm<sup>2</sup> to prepare an aerosol type hair dye.

As the propellant, there may be used a compressed gas such as nitrogen and laughing gas as well as carbon dioxide and a liquefied gas such as Furon 11, 12, 114 and LPG either alone or in combination.

This hair dye is coated on a white hair, which is treated at 30°C for 30 minutes, washed with water, shampooed and dried.

The white hair was dyed in grayish color.

This aerosol type hair dye has improved stability of PROD since it is brought into contact with air and/or the propellant during storage. Further, it is also possible to replace the piston can by a bag-in can and an EXXEL SYSTEM (product of CONTAINER INDUSTRIES INC., America) which ejects the contents by directly applying the rubber pressure to the container.

Example 7 Two-packages B	Hair D	ve (Shampoo	Type)
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Paraphenylenediamine	0.28%
Paraaminophenol	0.1
Orthoaminophenol	. 0.14
Nitroparaphenylenediamine	0.02
Resorcin	0.4
Polyoxyethylene lauryl ether	
sodium sulfate (3 E.O.)	10.0
Coconut oil fatty acid amide	
propyldimethylaminoacetic acid betaine	4.0
Coconut oil fatty acid diethanolamide	5.0

D-Glucose	5.5%
Propylene glycol	1.0
GOD (2.4 units/mg)	4.15
Mutarotase (0.95 units/mg)	0.1
Peroxidase (112 units/mg)	0.0029
0.1 M phosphate buffer (pH 6.5)	Balance

Note: GOD and mutarotase in this formulation are added just before the hair dyeing treatment.

This composition is coated on a white hair, which is treated at 30°C for 30 minutes, washed with water and dried.

The white hair was dyed in slightly purple-tinted brown color.

Further, like the conventional shampoo, this composition is coated on a white hair at 30°C for 3 to 5 minutes, then immediately washed with water and dried. When this procedure was repeated on the white hair once a day for 10 days, it was dyed in slightly purple-tinted brown color.

Example 8	Two-packages	Hair	Dye
[Coloring	Base]		

Orthoaminophenol Resorcin Sodium laurylsulfate D-Glucose 1 Disodium editate Ascorbic acid Ammonium hydroxide, Purified water Bal (The pH is adjusted to 7.0 with ammonium hydroxid [Enzyme solution] PROD (1 unit/mg) 20. Peroxidase (112 units/mg) 0. Glycerin 3.	
Resorcin Sodium laurylsulfate D-Glucose 1 Disodium editate Ascorbic acid Ammonium hydroxide, Purified water Bal (The pH is adjusted to 7.0 with ammonium hydroxid [Enzyme solution] PROD (1 unit/mg) 20. Peroxidase (112 units/mg) 0. Glycerin 3.	2.7%
Sodium laurylsulfate D-Glucose Disodium editate Ascorbic acid Ammonium hydroxide, Purified water Bal (The pH is adjusted to 7.0 with ammonium hydroxid [Enzyme solution] PROD (1 unit/mg) Peroxidase (112 units/mg) Glycerin 3.	0.2
D-Glucose 1 Disodium editate Ascorbic acid Ammonium hydroxide, Purified water Bal (The pH is adjusted to 7.0 with ammonium hydroxid [Enzyme solution] PROD (1 unit/mg) 20. Peroxidase (112 units/mg) 0. Glycerin 3.	0.5
Disodium editate Ascorbic acid Ammonium hydroxide, Purified water  (The pH is adjusted to 7.0 with ammonium hydroxid  [Enzyme solution]  PROD (1 unit/mg)  Peroxidase (112 units/mg)  Glycerin  20.	1.0
Ascorbic acid  Ammonium hydroxide, Purified water Bal  (The pH is adjusted to 7.0 with ammonium hydroxid  [Enzyme solution]  PROD (1 unit/mg) 20.  Peroxidase (112 units/mg) 0.  Glycerin 3.	11.0
Ammonium hydroxide, Purified water Bal (The pH is adjusted to 7.0 with ammonium hydroxid [Enzyme solution] PROD (1 unit/mg) Peroxidase (112 units/mg) Glycerin 3.	0.1
(The pH is adjusted to 7.0 with ammonium hydroxid [Enzyme solution]  PROD (1 unit/mg)  Peroxidase (112 units/mg)  Glycerin  20.	0.4
<pre>[Enzyme solution]    PROD (1 unit/mg) 20.    Peroxidase (112 units/mg) 0.    Glycerin 3.</pre>	alance
PROD (1 unit/mg) 20. Peroxidase (112 units/mg) 0. Glycerin 3.	ide.)
Peroxidase (112 units/mg) 0. Glycerin 3.	
Glycerin 3.	9.0%
•	0.006-
0 1M Phogphato buffer (PU C E)	3.0
0.1M Phosphate buffer (pH 6.5) Bal	alance
First, a solution obtained by mixing the coloring	ıg

base and the enzyme solution at a ratio of 1:1 is coated on a white hair, which is treated at 30°C for 30 minutes, then washed with water, shampooed and dried.

The white hair was dyed in grayish color.

## Example 9 Two-packages Hair Dye [Coloring Base]

Paraphenylenediamine	88.0
. Paraaminophenol	0.2
Orthoaminophenol	1.0
Resorcin	1.6
Oleic acid	20.0
Bis-2-Hydroxyethylsorbitanamine	9.0
Hydroxyethylstearylamide	6.0
Propylene glycol	12.0
Isopropanol	10.0
Disodium editate	0.3
Sodium sulfite	0.3
Ammonium hydroxide, purified water	Balance

(The pH is adjusted to 8.5 with ammonium hydroxide.) [Enxyme Powder]

Uricase (2.8 units/mg)		15.49
Uric acid	·	84.6

First, a solution obtained by mixing the coloring base and the enzyme powder at a ratio of 14:1 is coated on a white hair, which is treated at 30°C for 30 minutes, washed with water, shampooed, and dried.

The white hair was dyed in gray-tinted brown color.

#### Example 10 Three-packages Hair Dye

#### [Coloring Base]

Paraphenylenediamine	0.56์ร์
Paraaminophenol	0.20
Orthoaminophenol	0.28
Nitroparaphenylenediamine	0.04

Resorcin	0.80%
Liquid paraffin	5.0
Lanolin alcohol	2.0
Polyoxyethylene lauryl ether phosphate (3 E.O.)	2.0
Lauric acid diethanolamide	5:0
Thioglycolic acid	0.2
Disodium editate	0.1
0.1M Phosphate buffer (pH 6.5)	Balance
[Substrate Solution]	
D-Glucose	21.6%
Purified water	Balance
[Enzyme Solution]	
Mutarotase (0.95 unit/mg)	0.4%
GOD (2.4 units/mg)	16.6
Peroxidase (112 units/mg)	0.0116
Glycerin	3.0
<pre>0.lM Phosphate buffer (pH 7.0)</pre>	Balance

First, a solution obtained by mixing the coloring base, the substrate solution and the enzyme solution at a ratio of 2:1:1 is coated on a white hair, which is treated at 30°C for 30 minutes, washed with water, shampooed and dried.

The white hair was dyed in slightly purple-tinted brown color.

Example 11	Cold Waving Preparation Mainly Containing
	Thioglycolic Acid
	(Thioglycolic Acid-based Permanent Waving
	Agent)

[First Solution (Reducing Agent)]

Ammonium thioglycolate solution
(50% as thioglycolic acid)

Polyoxyethylene oleyl ether (10 E.O.)

" (20 E.O.)

Sodium lauryl sulfate

Hydrolyzed collagen solution (20%)

13.0%
1.0

0.5

Disodium editate

0.1%

Ammonium hydroxide, purified water

Balance

(The pH is adjusted to 9.1 with ammonium hydroxide.) [Second Solution (Oxidation Agent)]

PROD (1 unit/mg)	 20.0%
Peroxidase (112 units/mg)	0.0058
D-Glucose	3.6
Glycerin	3.0
Purified water	Balance

Note: PROD in the formulation is added just before the treatment with the second solution.

First, in a conventional manner, a hair is protected on the tip with paper, wound on a plastic rod of 1.5 cm in diameter, dipped in the first solution of the above-described composition at 30°C for 10 minutes, washed with running water for 1 minute, then dipped in the second solution at 30°C for 10 minutes, and washed with water, to thereby conduct permanent waving treatment.

As a result, the hair was imparted with a uniform wave from the root to the tip, and also the touch was good.

# Example 12 Cold Waving Preparation Mainly Containing Cysteine

(Cysteine-based Permanent Waving Agent)

[First Solution (Reducing Agent)]

L-Cysteine Hydrochloride	7.0%
Cetanol	0.5
Oleyl alcohol	. 0.5
Polyoxyethylene cetyl ether (10	) E.O.)
" (1:	5 E.O.) 1.0
Disodium editate	0.1
Monoethanolamine, purified wate	er Balance

(The pH is adjusted to 9.1 with monoethanolamine.) [Second Solution (Oxidation Agent)]

GOD (2.4 units/mg)

8.3%

Mutarotase (0.95 unit/mg)	0.2%
Peroxidase (112 units/mg)	0.0058
D-Glucose	3.6
Sorbitol	3.0
Purified water	Balance

Note: GOD in the formulation is added just before the treatment with the second solution.

Using the above-described composition, permanent waving treatment was conducted in a manner similar to that in Example 11.

As a result, the hair was imparted with a uniform wave from the root to the tip, and also the touch was good.

Example 13 Heat-Waving Preparation Mainly Containing Thioglycolic Acid

[First Solution (Reducing Agent)]

Ammon	ium	thioglyco	late
Humon	T GHI		

<b>▼</b> •		
(50% as thioglycolic acid)	.*	10.0%
Cetanol		0.5
Oleyl alchol		0.5
Polyoxyethyl cetyl ether (10 E.O.)		0.5
" (15 E.O.)	•	1.0
Disodium editate		0.1
Ammonium bicarbonate, purified water		Balance

(The pH is adjusted to 7.5 with ammonium bicarbonate.)

[Second Solution (Oxidation Agent)]

The same agent as the second solution in Example 11 is used.

Using the above-described composition, permanent waving treatment was conducted in a manner similar to that in Example 10, except that both treatments with the first solution and the second solution were conducted at 45°C.

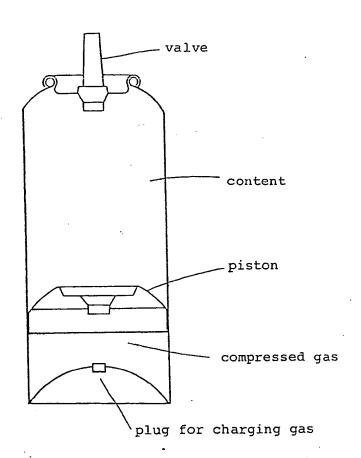
As a result, the hair was imparted with a uniform wave from the root to the tip, and also the touch was good.

#### SCOPE OF CLAIMS

- 1. A hair preparation composition characterized in that at least one dielectron reducing oxidase utilizing oxygen as an acceptor is incorporated.
- 2. The hair preparation composition according to Claim 1, wherein the dielectron reducing oxidase is at least one member selected from the group consisting of pyranose oxidase, glucose oxidase, glycerol oxidase, lactic acid oxidase, pyruvic acid oxidase and uricase.
- 3. A hair preparation composition which comprises an enzyme recited in Claim 1 or 2 and a donor of said enzyme.
- 4. A hair dye composition which consists of three packages, i.e. a composition comprising an oxidation dye, a composition comprising an enzyme recited in Claim 1 or 2 and a composition comprising a donor of said enzyme.
- 5. A hair dye composition which consists of two packages, i.e. a package comprising one component selected from the group consisting of an oxidation dye, an enzyme recited in Claim 1 or 2 and a donor of said enzyme; and a package comprising the remaining two components.
- 6. A one-package type hair dye composition which comprises an oxidation dye, an enzyme recited in Claim 1 or 2 and a donor of said enzyme.
- 7. An oxidation agent composition for permanent waving preparation which comprises an enzyme recited in Claim 1 or 2 and a donor of said enzyme.
- 8. An oxidation agent composition for permanent waving preparation which consists of two packages, i.e. a package comprising an enzyme recited in Claim 1 or 2 and a package comprising a donor of said enzyme.
- 9. A hair preparation composition which comprises an -- enzyme recited in Claim 1 or 2, a donor of said enzyme and a peroxidase.
  - 10. The hair preparation composition according to Claim

1 or 2, wherein the oxidase is glucose oxidase and mutarotase is further contained.

Fig. 1



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## international search report

International Application No

PCT/JP88/00315

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х	US, A, 3,251,742 (Revlon, 17 March 1966 (17. 03. 66) Claim	Inc.)	1, 3-6, 9
x	JP, A, 47-10400 (The Procto 25 May 1972 (25. 05. 72) Claim & DE, A, 2155390 & FR, A, 2112550 & GB, A,		1, 3-6, 9
X	JP, A, 50-95436 (Tanabe Se 29 July 1975 (29. 07. 75) 712im (Family, Done) 2 DE, A, 2545986 & IR, A,		1, 3, 7-9
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Jun	e 18, 1988 (18. 06. 88)	July 4, 1988 (04.	07. 88)
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Japanese Patent Office			